This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1. (CURRENTLY AMENDED) An organic azide compound having the formula:

wherein Ar is an aromatic or a heteroaromatic radical derived from the group consisting of benzenes, polyfluorobenzenes, naphthalenes, naphthoquinones, anthracenes, anthraquinones, phenanthrenes, tetracenes, naphthacenediones, pyridines, quinolines, isoquinolines, indoles, isoindoles, pyrroles, imidiazoles, pyrazoles, pyrazines, purines, benzimidazoles, benzofurans, dibenzofurans, carbazoles, acridines, acridones, phenanthridines, thiophenes, benzothiophenes, dibenzothiophenes, xanthenes, xanthones, flavones, coumarins, and anthacylines;

E is selected from the group consisting of hydrogen-somatostatin receptor binding molecules, ST receptor binding molecules, neurotensin receptor binding molecules, bombesin receptor binding molecules, CCK receptor binding molecules, steroid receptor binding molecules, and carbohydrate receptor binding molecules;

L is selected from the group consisting of -(CH₂)_a-, -(CH₂)_bCONR¹-, -N(R²)CO(CH₂)_c-, -OCO(CH₂)_o-, -(CH₂)_eCO₂-, -OCONH-, -OCO₂-, -HNCONH-, -HNCSNH-, -HNNHCO-, -OSO₂-, -NR³(CH₂)_eCONR⁴-, -CONR⁵(CH₂)_fNR⁶CO-, and -NR⁷CO(CH₂)_gCONR⁸-;

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X is either a single bond or is selected from the group consisting of $-(CH_2)_h$, -OCO, -HNCO, $-(CH_2)_lCO$, and $-(CH_2)_lOCO$;

R¹ to R⁸ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, -OH, C1-C10 polyhydroxyalkyl, C1-C10 alkoxyl, C1-C10 alkoxyl, C1-C10 alkoxyl, C1-C10 alkoxyl, -SO₃H, -(CH₂)_kCO₂H, and -(CH₂)_lNR⁹R¹⁰;

R⁹ and R¹⁰ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, C5-C10 aryl, and C1-C10 polyhydroxyalkyl; and a to I independently range from 0 to 10.

2. (WITHDRAWN) The compound of claim 1 wherein Ar is an aromatic or heteroaromatic radical derived from polyfluorbenzenes; E is selected from the group consisting of somatostatin receptor binding molecules, ST receptor binding molecules, neurotensin receptor binding molecules, bombesin receptor binding molecules, CCK receptor binding molecules, and steroid receptor binding molecules; L is selected from the group consisting of -(CH₂)₆CONR¹-, -N(R²)CO(CH₂)₆-, -OCO(CH₂)₆-, -(CH₂)₆CO₂-, -HNCONH-, -HNCSNH-, and -NR⁷CO(CH₂)₆CONR⁸-; X is either a single bond or is selected from the group consisting of -(CH₂)_h-, -OCO-, -(CH₂)_iCO-, and -(CH₂)_iOCO-, R¹, R², R⁷ and R⁸ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, C1-C10 polyhydroxyalkyl, -(CH₂)_kCO₂H, and -(CH₂)_iNR⁹R¹⁰; R⁹ and R¹⁰ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, and C1-C10 polyhydroxyalkyl; and subscripts b-e and g-j independently range from 0 to 6.

- 3. (WITHDRAWN) The compound of claim 1 wherein Ar is an aromatic or heteroaromatic radical derived from anthraquinones; E is selected from the group consisting of somatostatin receptor binding molecules, ST receptor binding molecules, neurotensin receptor binding molecules, bombesin receptor binding molecules, CCK receptor binding molecules, and steroid receptor binding molecules; L is selected from the group consisting of -(CH₂)₀CONR¹-, -N(R²)CO(CH₂)_c-, -OCO(CH₂)_d-, -(CH₂)_eCO₂-, -HNCONH-, -HNCSNH-, and -NR⁷CO(CH₂)₀CONR⁸-; X is either a single bond or is selected from the group consisting of -(CH₂)_h-, -OCO-, -(CH₂)_hCO-, and -(CH₂)_hOCO-., R¹, R², R⁷ and R⁸ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, C1-C10 polyhydroxyalkyl, -(CH₂)_kCO₂H, and -(CH₂)_hNR⁹R¹⁰; R⁹ and R¹⁰ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, and C1-C10 polyhydroxyalkyl; and subscripts b-e and g-j independently range from 0 to 6.
- 4. (WITHDRAWN) The compound of claim 1 wherein Ar is an aromatic or heteroaromatic radical derived from napthacenediones; E is selected from the group consisting of somatostatin receptor binding molecules, ST receptor binding molecules, neurotensin receptor binding molecules, bombesin receptor binding molecules, CCK receptor binding molecules, and steroid receptor binding molecules; L is selected from the group consisting of -(CH₂)₆CONR¹-, -N(R²)CO(CH₂)₆-, -OCO(CH₂)₆-, -(CH₂)₆CO₂-, -HNCONH-, -HNCSNH-, and -NR⁷CO(CH₂)₉CONR⁸-; X is either a single bond or is selected from the group consisting of -(CH₂)₆-, -OCO-, -(CH₂)₆CO-, and -(CH₂)₇OCO-., R¹, R², R⁷ and R⁸ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, C1-C10 polyhydroxyalkyl, -(CH₂)₆CO₂H, and -(CH₂)₁NR⁹R¹⁰; R⁹ and R¹⁰

are independently selected from the group consisting of hydrogen, C1-C10 alkyl, and C1-C10 polyhydroxyalkyl; and subscripts b-e and g-j independently range from 0 to 6.

- 5. (WITHDRAWN) The compound of claim 1 wherein Ar is an aromatic or heteroaromatic radical derived from indoles; E is selected from the group consisting of somatostatin receptor binding molecules, ST receptor binding molecules, neurotensin receptor binding molecules, bombesin receptor binding molecules, CCK receptor binding molecules, and steroid receptor binding molecules; L is selected from the group consisting of -(CH₂)₆CONR¹-, -N(R²)CO(CH₂)₆-, -OCO(CH₂)₆-, -(CH₂)₆CO₂-, -HNCONH-, -HNCSNH-, and -NR⁷CO(CH₂)₆CONR⁸-; X is either a single bond or is selected from the group consisting of -(CH₂)₆-, -OCO-, -(CH₂)₆CO-, and -(CH₂)₆OCO-., R¹, R², R⁷ and R⁸ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, C1-C10 polyhydroxyalkyl, -(CH₂)₆CO₂H, and -(CH₂)₆NR⁹R¹⁰; R⁹ and R¹⁰ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, and C1-C10 polyhydroxyalkyl; and subscripts b-e and g-j independently range from 0 to 6.
- 6. (WITHDRAWN) The compound of claim 1 wherein Ar is an aromatic or heteroaromatic radical derived from acridines; E is selected from the group consisting of somatostatin receptor binding molecules, ST receptor binding molecules, neurotensin receptor binding molecules, bombesin receptor binding molecules, CCK receptor binding molecules, and steroid receptor binding molecules; L is selected from the group consisting of -(CH₂)₆CONR¹-, -N(R²)CO(CH₂)₆-, -OCO(CH₂)₆-, -(CH₂)₆CO₂-, -HNCONH-, -HNCSNH-, and -NR⁷CO(CH₂)₉CONR⁸-; X is either a single bond or is Page 5 of 24

selected from the group consisting of -(CH₂)_h-, -OCO-, -(CH₂)_iCO-, and -(CH₂)_jOCO-., R¹, R², R⁷ and R⁸ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, C1-C10 polyhydroxyalkyl, -(CH₂)_kCO₂H, and -(CH₂)_iNR⁹R¹⁰; R⁹ and R¹⁰ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, and C1-C10 polyhydroxyalkyl; and subscripts b-e and g-j independently range from 0 to 6.

- 7. (WITHDRAWN) The compound of claim 1 wherein Ar is an aromatic or heteroaromatic radical derived from acridones; E is selected from the group consisting of somatostatin receptor binding molecules, ST receptor binding molecules, neurotensin receptor binding molecules, bombesin receptor binding molecules, CCK receptor binding molecules, and steroid receptor binding molecules; L is selected from the group consisting of -(CH₂)₆CONR¹-, -N(R²)CO(CH₂)₆-, -OCO(CH₂)₆-, -(CH₂)₆CO₂-, -HNCONH-, -HNCSNH-, and -NR⁷CO(CH₂)₆CONR⁹-; X is either a single bond or is selected from the group consisting of -(CH₂)₆CONR⁹-; X is either a single bond or is selected from the group consisting of -(CH₂)₆CO₂-, -(CH₂)₆CO₂-, and -(CH₂)₆OCO₋. R¹, R², R⁷ and R⁸ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, C1-C10 polyhydroxyalkyl, -(CH₂)₆CO₂H, and -(CH₂)₆NR⁹R¹⁰; R⁹ and R¹⁰ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, and C1-C10 polyhydroxyalkyl; and subscripts b-e and g-j independently range from 0 to 6.
- 8. (CURRENTLY AMENDED) The compound of claim 1 wherein Ar is an aromatic or a heteroaromatic radical derived from phenanthridines; E is selected from the group Page 6 of 24

consisting of somatostatin receptor binding molecules, ST receptor binding molecules, neurotensin receptor binding molecules, bombesin receptor binding molecules, CCK receptor binding molecules, and steroid receptor binding molecules; L is selected from the group consisting of -(CH₂)_bCONR¹-, -N(R²)CO(CH₂)_c-, -OCO(CH₂)_o-, -(CH₂)_eCO₂-, -HNCONH-, -HNCSNH-, and -NR⁷CO(CH₂)_gCONR⁸-; X is either a single bond or is selected from the group consisting of -(CH₂)_h-, -OCO-, -(CH₂)_tCO-, and -(CH₂)_tOCO-, R¹, R², R⁷ and R⁸ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, C1-C10 polyhydroxyalkyl, -(CH₂)_kCO₂H, and -(CH₂)_tNR⁹R¹⁰; R⁹ and R¹⁰ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, and C1-C10 polyhydroxyalkyl; and subscripts b-e and g-j independently range from 0 to 6.

9. (WITHDRAWN) The compound of claim 1 wherein Ar is an aromatic or heteroaromatic radical derived from xanthones; E is selected from the group consisting of somatostatin receptor binding molecules, ST receptor binding molecules, neurotensin receptor binding molecules, bombesin receptor binding molecules, CCK receptor binding molecules, and steroid receptor binding molecules; L is selected from the group consisting of -(CH₂)_bCONR¹-, -N(R²)CO(CH₂)_c-, -OCO(CH₂)_d-, -(CH₂)_eCO₂-, -HNCONH-, -HNCSNH-, and -NR⁷CO(CH₂)_gCONR⁸-; X is either a single bond or is selected from the group consisting of -(CH₂)_h-, -OCO-, -(CH₂)_iCO-, and -(CH₂)_jOCO-., R¹, R², R⁷ and R⁹ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, C1-C10 polyhydroxyalkyl, -(CH₂)_kCO₂H, and -(CH₂)_iNR⁹R¹⁰; R⁹ and R¹⁰ are independently selected from the group consisting of hydrogen,

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C1-C10 alkyl, and C1-C10 polyhydroxyalkyl; and subscripts b-e and g-j independently range from 0 to 6.

10. (WITHDRAWN) The compound of claim 1 wherein Ar is an aromatic or heteroaromatic radical derived from anthracyclines; E is selected from the group consisting of somatostatin receptor binding molecules, ST receptor binding molecules, neurotensin receptor binding molecules, bombesin receptor binding molecules, CCK receptor binding molecules, and steroid receptor binding molecules; L is selected from the group consisting of -(CH₂)₆CONR¹-, -N(R²)CO(CH₂)₆-, -OCO(CH₂)₆-, -(CH₂)₆CO₂-, -HNCONH-, -HNCSNH-, and -NR⁷CO(CH₂)₉CONR⁸-; X is either a single bond or is selected from the group consisting of -(CH₂)_h-, -OCO-, -(CH₂)_lCO-, and -(CH₂)_lOCO-., R¹, R², R⁷ and R⁸ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, C1-C10 polyhydroxyalkyl, -(CH₂)_kCO₂H, and -(CH₂)_kNR⁹R¹⁰; R⁹ and R¹⁰ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, and C1-C10 polyhydroxyalkyl; and subscripts b-e and g-j independently range from 0 to 6.

11. (CURRENTLY AMENDED) A method of performing a phototherapeutic procedure which comprises:

(a) administering to a target tissue in an animal an effective amount of an organic azide photosensitizer having the formula

wherein Ar is an aromatic or a heteroaromatic radical derived from the group consisting of benzenes, polyfluorobenzenes, naphthalenes, naphthoquinones, anthracenes, anthraquinones, phenanthrenes, tetracenes, naphthacenediones, pyridines, quinolines, isoquinolines, indoles, isoindoles, pyrroles, imidiazoles, pyrazoles, pyrazines, purines, benzimidazoles, benzofurans, dibenzofurans, carbazoles, acridines, acridones, phenanthridines, thiophenes, benzothiophenes, dibenzothiophenes, xanthenes, xanthones, flavones, coumarins, and anthacylines; E is a hydrogen atom or is selected from the group consisting of somatostatin receptor binding molecules, ST receptor binding molecules, neurotensin receptor binding molecules, bombesin receptor binding molecules, CCK receptor binding molecules, steroid receptor binding molecules, and carbohydrate receptor binding molecules; L is selected from the group consisting of $-(CH_2)_a$ -, $-(CH_2)_bCONR^1$ -, $-N(R^2)CO(CH_2)_c$ -, $-OCO(CH_2)_d$ -, $-(CH_2)_aCO_2$ -, $-OCONH_2$, -OCO₂-, -HNCONH-, -HNCSNH-, -HNNHCO-, -OSO₂-, -NR³(CH₂), CONR⁴-, -CONR⁵(CH₂),NR⁶CO-, and -NR⁷CO(CH₂),CONR⁶-; X is either a single bond or is selected from the group consisting of -(CH₂)_h-, -OCO-, -HNCO-, -(CH₂)_iCO-, and -(CH₂)_iOCO-; R¹ to R⁸ are independently selected from the group consisting of

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procedure.

hydrogen, C1-C10 alkyl, -OH, C1-C10 polyhydroxyalkyl, C1-C10 alkoxyl, C1-C10 alkyl, C2-C10 aryl, and C1-C10 polyhydroxyalkyl; and subscripts a to I independently range from 0 to 10; and (b) exposing said target tissues with the light of wavelength between 300 and 950 nm with sufficient power and fluence rate to perform the phototherapeutic

12. (WITHDRAWN) The method of claim 11, wherein Ar is an aromatic or heteroaromatic radical derived from polyfluorbenzenes; E is selected from the group consisting of somatostatin receptor binding molecules, ST receptor binding molecules, neurotensin receptor binding molecules, bombesin receptor binding molecules, CCK receptor binding molecules, and steroid receptor binding molecules; L is selected from the group consisting of -(CH₂)_bCONR¹-, -N(R²)CO(CH₂)_c-, -OCO(CH₂)_d-, -(CH₂)_eCO₂-, -HNCONH-, -HNCSNH-, and -NR⁷CO(CH₂)_gCONR⁶-; X is either a single bond or is selected from the group consisting of -(CH₂)_h-, -OCO-, -(CH₂)_hCO-, and -(CH₂)_iOCO-., R¹, R², R⁷ and R⁶ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, C1-C10 polyhydroxyalkyl, -(CH₂)_kCO₂H, and -(CH₂)_iNR⁹R¹⁰; R⁹ and R¹⁰ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, and C1-C10 polyhydroxyalkyl; and subscripts b-e and g-j independently range from 0 to 6.

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- 13. (WITHDRAWN) The method of claim 11, wherein Ar is an aromatic or heteroaromatic radical derived from anthraquinones; E is selected from the group consisting of somatostatin receptor binding molecules, ST receptor binding molecules, neurotensin receptor binding molecules, bombesin receptor binding molecules, CCK receptor binding molecules, and steroid receptor binding molecules; L is selected from the group consisting of -(CH₂)₆CONR¹-, -N(R²)CO(CH₂)₆-, -OCO(CH₂)₆-, -(CH₂)₆CO₂-, -HNCONH-, -HNCSNH-, and -NR⁷CO(CH₂)₉CONR⁸-; X is either a single bond or is selected from the group consisting of -(CH₂)_h-, -OCO-, -(CH₂)_hCO-, and -(CH₂)_hOCO-., R¹, R², R⁷ and R⁸ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, C1-C10 polyhydroxyalkyl, -(CH₂)_kCO₂H, and -(CH₂)_hNR⁹R¹⁰; R⁹ and R¹⁰ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, and C1-C10 polyhydroxyalkyl; and subscripts b-e and g-j independently range from 0 to 6.
- 14. (WITHDRAWN) The method of claim 11, wherein Ar is an aromatic or heteroaromatic radical derived from napthacenediones; E is selected from the group consisting of somatostatin receptor binding molecules, ST receptor binding molecules, neurotensin receptor binding molecules, bombesin receptor binding molecules, CCK receptor binding molecules, and steroid receptor binding molecules; L is selected from the group consisting of $-(CH_2)_bCONR^1$ -, $-N(R^2)CO(CH_2)_c$ -, $-OCO(CH_2)_d$ -, $-(CH_2)_eCO_2$ -, -HNCONH-, -HNCSNH-, and $-NR^7CO(CH_2)_gCONR^8$ -; X is either a single bond or is selected from the group consisting of $-(CH_2)_h$ -, -OCO-, $-(CH_2)_iCO$ -, and $-(CH_2)_jOCO$ -., R^1 , R^2 , R^7 and R^8 are independently selected from the group consisting of hydrogen, C1-C10 alkyl, C1-C10 polyhydroxyalkyl, $-(CH_2)_kCO_2H$, and $-(CH_2)_iNR^8R^{10}$; R^9 and R^{10}

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are independently selected from the group consisting of hydrogen, C1-C10 alkyl, and C1-C10 polyhydroxyalkyl; and subscripts b-e and g-j independently range from 0 to 6.

15. (WITHDRAWN) The method of claim 11, wherein Ar is an aromatic or heteroaromatic radical derived from indoles; E is selected from the group consisting of somatostatin receptor binding molecules, ST receptor binding molecules, neurotensin receptor binding molecules, bombesin receptor binding molecules, CCK receptor binding molecules, and steroid receptor binding molecules; L is selected from the group consisting of -(CH₂)_bCONR¹-, -N(R²)CO(CH₂)_c-, -OCO(CH₂)_d-, -(CH₂)_aCO₂-, -HNCONH-, -HNCSNH-, and -NR⁷CO(CH₂)_bCONR⁸-; X is either a single bond or is selected from the group consisting of -(CH₂)_n-, -OCO-, -(CH₂)_iCO-, and -(CH₂)_iOCO-., R¹, R², R⁷ and R⁸ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, C1-C10 polyhydroxyalkyl, -(CH₂)_kCO₂H, and -(CH₂)_iNR⁹R¹⁰; R⁹ and R¹⁰ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, and C1-C10 polyhydroxyalkyl; and subscripts b-e and g-j independently range from 0 to 6.

16. (WITHDRAWN) The method of claim 11, wherein Ar is an aromatic or heteroaromatic radical derived from acridines; E is selected from the group consisting of somatostatin receptor binding molecules, ST receptor binding molecules, neurotensin receptor binding molecules, bombesin receptor binding molecules, CCK receptor binding molecules, and steroid receptor binding molecules; L is selected from the group consisting of -(CH₂)_bCONR¹-, -N(R²)CO(CH₂)_c-, -OCO(CH₂)_d-, -(CH₂)_eCO₂-, -HNCONH-, -HNCSNH-, and -NR²CO(CH₂)_gCONR³-; X is either a single bond or is

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selected from the group 1 consisting of -(CH₂)_h-, -OCO-, -(CH₂)_iCO-, and -(CH₂)_jOCO-., R¹, R², R⁷ and R⁸ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, C1-C10 polyhydroxyalkyl, -(CH₂)_kCO₂H, and -(CH₂)_iNR⁹R¹⁰; R⁹ and R¹⁰ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, and C1-C10 polyhydroxyalkyl; and subscripts b-e and g-j independently range from 0 to 6.

- 17. (WITHDRAWN) The method of claim 11, wherein Ar is an aromatic or heteroaromatic radical derived from acridones; E is selected from the group consisting of somatostatin receptor binding molecules, ST receptor binding molecules, neurotensin receptor binding molecules, bombesin receptor binding molecules, CCK receptor binding molecules, and steroid receptor binding molecules; L is selected from the group consisting of -(CH₂)_bCONR¹-, -N(R²)CO(CH₂)_c-, -OCO(CH₂)_d-, -(CH₂)_eCO₂-, -HNCONH-, -HNCSNH-, and -NR⁷CO(CH₂)_gCONR⁸-; X is either a single bond or is selected from the group consisting of -(CH₂)_b-, -OCO-, -(CH₂)_cCO-, and -(CH₂)_fOCO-., R¹, R², R⁷ and R⁸ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, C1-C10 polyhydroxyalkyl, -(CH₂)_kCO₂H, and -(CH₂)_iNR⁹R¹⁰; R⁹ and R¹⁰ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, and C1-C10 polyhydroxyalkyl; and subscripts b-e and g-j independently range from 0 to 6.
- 18. (WITHDRAWN) The method of claim 11, wherein Ar is an aromatic or heteroaromatic radical derived from phenanthridines; E is selected from the group

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consisting of somatostatin receptor binding molecules, ST receptor binding molecules, neurotensin receptor binding molecules, bombesin receptor binding molecules, CCK receptor binding molecules, and steroid receptor binding molecules; L is selected from the group consisting of - $(CH_2)_bCONR^1$ -, - $N(R^2)CO(CH_2)_c$ -, - $OCO(CH_2)_d$ -, - $(CH_2)_bCO_2$ -, -HNCONH-, -HNCSNH-, and - $NR^7CO(CH_2)_gCONR^8$ -; X is either a single bond or is selected from the group consisting of - $(CH_2)_h$ -, -OCO-, - $(CH_2)_lCO$ -, and - $(CH_2)_lOCO$ -., R^1 , R^2 , R^7 and R^8 are independently selected from the group consisting of hydrogen, C1-C10 alkyl, C1-C10 polyhydroxyalkyl, - $(CH_2)_kCO_2H$, and - $(CH_2)_lNR^9R^{10}$; R^9 and R^{10} are independently selected from the group consisting of hydrogen, C1-C10 alkyl, and C1-C10 polyhydroxyalkyl; and subscripts b-e and g-j independently range from 0 to 6.

19. (WITHDRAWN) The method of claim 11, wherein Ar is an aromatic or heteroaromatic radical derived from xanthones; E is selected from the group consisting of somatostatin receptor binding molecules, ST receptor binding molecules, neurotensin receptor binding molecules, bombesin receptor binding molecules, CCK receptor binding molecules, and steroid receptor binding molecules; L is selected from the group consisting of -(CH₂)₆CONR¹-, -N(R²)CO(CH₂)_c-, -OCO(CH₂)_d-, -(CH₂)_eCO₂-, -HNCONH-, -HNCSNH-, and -NR⁷CO(CH₂)_eCONR⁸-; X is either a single bond or is selected from the group consisting of -(CH₂)_h-, -OCO-, -(CH₂)_iCO-, and -(CH₂)_iOCO-., R¹, R², R⁷ and R⁸ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, C1-C10 polyhydroxyalkyl, -(CH₂)_kCO₂H, and -(CH₂)_iNR⁹R¹⁰; R⁹

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and R¹⁰ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, and C1-C10 polyhydroxyalkyl; and subscripts b-e and g-j independently range from 0 to 6.

- 20. (WITHDRAWN) The method of clalm 11, wherein Ar is an aromatic or heteroaromatic radical derived from anthracyclines; E is selected from the group consisting of somatostatin receptor binding molecules, ST receptor binding molecules, neurotensin receptor binding molecules, bombesin receptor binding molecules, CCK receptor binding molecules, and steroid receptor binding molecules; L is selected from the group consisting of -(CH₂)₆CONR¹-, -N(R²)CO(CH₂)₆-, -OCO(CH₂)₆-, -(CH₂)₆CO₂-, -HNCONH-, -HNCSNH-, and -NR⁷CO(CH₂)₆CONR⁸-; X is either a single bond or is selected from the group consisting of -(CH₂)₆-, -OCO-, -(CH₂)₆CO-, and -(CH₂)₆OCO-... R¹, R², R⁷ and R⁸ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, C1-C10 polyhydroxyalkyl, -(CH₂)₆CO₂H, and -(CH₂)₁NR⁹R¹⁰; R⁹ and R¹⁰ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, and C1-C10 polyhydroxyalkyl; and subscripts b-e and g-j independently range from 0 to 6.
- 21. (WITHDRAWN) The method of claim 11 further comprising the step of allowing said photosensitizer to accumulate in said target tissue before exposing said tissue to light.
- 22. (WITHDRAWN) The method of claim 11 wherein the photosensitizer is in a concentration ranging from about 1 nM to about 0.5 M.

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- 23. (WITHDRAWN) The method of claim 11 wherein the photosensitizer is in a concentration ranging from 1 μM to 10 mM.
- 24. (WITHDRAWN) The method of claim 11 wherein the photosensitizer is parenterally administered within a formulation including pharmaceutically acceptable substances selected from the group consisting of buffers, emulsifiers, surfactants, electrolytes, and combinations thereof.
- 25. (WITHDRAWN) The method of claim 11 wherein the photosensitizer is administered by a method selected from the group consisting of aerosol spray, cutaneously, parenterally, enterally, and topically.
- 26. (WITHDRAWN) The method of claim 11 wherein the effective amount of the photosensitizer administered is in the range of 0.1 mg/kg body weight to 500 mg/kg body weight.
- 27. (WITHDRAWN) The method of claim 11 wherein the effective amount of the photosensitizer administered is in the range of 0.5 mg/kg body weight to 2 mg/kg body weight.
- 28. (NEW) A composition comprising an organic azide compound having the formula:

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wherein Ar is a heteroaromatic phenanthrene radical;

E is a somatostatin receptor binding molecule,

L is selected from the group consisting of -(CH₂)_a-, -(CH₂)_bCONR¹-, -N(R²)CO(CH₂)_c-, -OCO(CH₂)_d-, -(CH₂)_eCO₂-, -OCONH-, -OCO₂-, -HNCONH-, -HNCSNH-, -HNNHCO-, -OSO₂-, -NR³(CH₂)_eCONR⁴-, -CONR⁵(CH₂)_tNR⁶CO-, and -NR⁷CO(CH₂)_gCONR⁸-;

X is either a single bond or is selected from the group consisting of $-(CH_2)_h$ -, -OCO-, -HNCO-, $-(CH_2)_lCO$ -, and $-(CH_2)_jOCO$ -;

R¹ to R⁸ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, -OH, C1-C10 polyhydroxyalkyl, C1-C10 alkoxyl, C1-C10 alkoxyl, C1-C10 alkoxyl, C1-C10 alkoxyl, -SO₃H, -(CH₂)_kCO₂H, and -(CH₂)_lNR⁹R¹⁰;

R⁹ and R¹⁰ are independently selected from the group consisting of hydrogen, C1-C10 alkyl, C5-C10 aryl, and C1-C10 polyhydroxyalkyl; and

a to I independently range from 0 to 10 in a pharmaceutically accepable formulation.